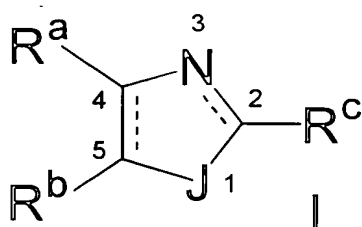
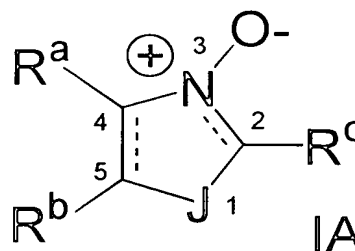


What is claimed:

1. A method of decreasing intraocular pressure or improving ocular accommodation in an animal, including a human, comprising administering an intraocular pressure decreasing amount or ocular accommodation improving amount of a compound of the formula I or IA,



or



wherein:

- a. J is oxygen, sulfur, or N-R^d;
 - 10 b. the carbon 2 to nitrogen bond is a double bond except when R^c is oxo;
 - c. the bond between carbons 4 and 5 is a single bond or a double bond;
 - d. R^a and R^b are
2. independently selected from hydrogen, acylamino, acyloxyalkyl, alkanoyl, alkanoylalkyl, alkenyl, alkoxy, alkoxycarbonyl, alkoxycarbonylalkyl, alkyl, alkylamino, (C₁-C₃)alkylenedioxy, allyl, amino, ω-alkylenesulfonic acid, carbamoyl, carboxy, carboxyalkyl (which alkyl can be substituted with alkyloxyimino), cycloalkyl, dialkylamino, halo, hydroxy, (C₂-C₆)hydroxyalkyl, mercapto, nitro, sulfamoyl, sulfonic acid, alkylsulfonyl, alkylsulfinyl, alkylthio, trifluoromethyl, morpholin-4-yl, thiomorpholin-4-yl, piperidin-1-yl, 4-[C₆ or C₁₀] arylpiperidin-1-yl, 4-[C₆ or C₁₀] arylpiperazin-1-yl, Ar {wherein, consistent with the rules of aromaticity, Ar is C₆ or C₁₀ aryl or a 5- or 6-membered heteroaryl ring, wherein the 6-membered heteroaryl ring contains one to three atoms of N, and the 5-membered heteroaryl ring contains from one to three atoms of N or one atom of O or S and zero to two atoms of N, each heteroaryl ring can be fused to a substituted benzene, pyridine, pyrimidine, pyrazine, pyridazine, or (1,2,3)triazine (wherein the ring fusion is at a carbon-carbon double bond of Ar)}, Ar-alkyl, ArO-, ArSO₂-, ArSO-, ArS-, ArSO₂NH-, ArNH-, (N-Ar)(N-alkyl)N-, ArC(O)-,

ArC(O)NH-, ArNH-C(O)-, and (N-Ar)(N-alkyl)N-C(O)-, or together R^a and R^b comprise methylenedioxy-; or

2. together with their ring carbons form a C₆- or C₁₀- aryl fused ring; or
3. together with their ring carbons form a C₅-C₇ fused cycloalkyl ring having up to
 5 two double bonds including any fused double bond of the containing group,
 which cycloalkyl ring can be substituted by one or more of the group consisting
 of alkyl, alkoxycarbonyl, amino, aminocarbonyl, carboxy, fluoro, or oxo; or
4. together with their ring carbons form a fused 5- or 6-membered heteroaryl ring,
 wherein the 6-membered heteroaryl ring contains one to three atoms of N, and
 10 the 5-membered heteroaryl ring contains from one to three atoms of N or one
 atom of O or S and zero to two atoms of N; or
5. together with their ring carbons form a fused five to eight membered second
 heterocycle, wherein the fused heterocycle consists of ring atoms selected from
 the group consisting of carbon, nitrogen, oxygen, sulfur, and S(O)_n, wherein n is
 15 1 or 2;

e. R^d is alkyl, alkenyl, hydrogen, or Ar;

f. R^c is

1. oxo (when $\Delta^{2,3}$ is not present), or (when $\Delta^{2,3}$ is present) hydrogen, alkyl, alkylthio,
 hydrogen, ^{thiol}mercapto, amino, amino(C₁-C₅)alkyl, or amino(C₆ or C₁₀)aryl, or
 20 wherein the amino of the last three groups can be substituted with

(a) Ar,

(b) Ar-Z-, Ar-alkyl-Z-, Ar-Z-alkyl, Ar-amino-Z-, Ar-aminoalkyl-Z-, or
 Ar-oxyalkyl-Z-, wherein Z is a carbonyl or -SO₂-

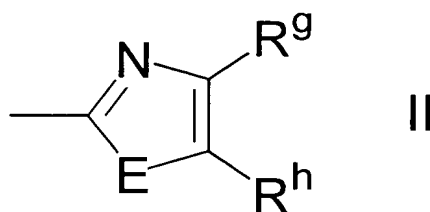
(c) formyl or alkanoyl,

2. -NHC(O)(CH₂)_n-D-R^eR^f, wherein D is oxygen, sulfur or nitrogen, wherein where
 25 D is nitrogen n is 0, 1 or 2, but when D is oxygen or sulfur n=1 or 2, and R^f is
 present only when D is nitrogen,
 wherein

(a) R^e is

(1) Ar,

(2) a group of the formula



wherein E is sulfur, oxygen, or N-Rⁱ, and R^g, R^h and Rⁱ are independently the same as R^a, R^b and R^d, respectively,

- 5 **(3)** a C₃-C₈ cycloalkyl ring having up to one double bond with the proviso that the carbon linking the cycloalkyl ring to D is saturated, which cycloalkyl ring can be substituted by one or more alkyl-, alkoxy carbonyl-, amino-, aminocarbonyl-, carboxy-, fluoro-, or oxo-substituents;
- 10 **(4)** a 5- or 6-membered heteroaryl ring containing at least one and up to three atoms of N for the 6-membered heteroaryl rings and from one to three atoms of N or one atom of O or S and zero to two atoms of N for the 5-membered heteroaryl rings;
- 15 **(5)** hydrogen, (C₂-C₆)hydroxyalkyl, alkanoylalkyl, alkyl, alkoxy carbonylalkyl, alkenyl, carboxyalkyl (which alkyl can be substituted with alkoxyimino), alkoxy carbonyl, a group Ar^ϕ which is C₆- or C₁₀- aryl or a 5- or 6-membered, or 9- or 10-membered heteroaryl (wherein the heteroatom is one oxygen, one sulfur or one nitrogen) or Ar^ϕ-alkyl; and
- 20 **(b)** R^f is independently hydrogen, (C₂-C₆)hydroxyalkyl, alkanoylalkyl, alkyl, alkoxy carbonylalkyl, alkenyl, carboxyalkyl (which alkyl can be substituted with alkoxyimino), alkoxy carbonyl, Ar^ϕ, or Ar^ϕ-alkyl;

wherein aryl, Ar, or Ar^ϕ can be substituted with, in addition to any substitutions specifically noted one or more substituents selected from the group of acylamino, acyloxyalkyl, alkanoyl, alkanoylalkyl, alkenyl, alkoxy, alkoxy carbonyl, alkoxy carbonylalkyl, alkyl, alkylamino, (C₁-C₃)alkylenedioxy, alkylsulfonyl, alkylsulfinyl, ω-alkylenesulfonic acid, alkylthio, allyl, amino, ArC(O)-, ArC(O)NH-, carboxy, carboxyalkyl, cycloalkyl, dialkylamino, halo, trifluoromethyl, hydroxy, (C₂-C₆)hydroxyalkyl, mercapto, nitro, ArO-, Ar-, Ar-

alkyl-, sulfamoyl, sulfonic acid, 1-pyrrolidinyl, 4-[C₆ or C₁₀]arylpiperazin-1-yl-, 4-[C₆ or C₁₀]arylpiperidin-1-yl, azetidin-1-yl, morpholin-4-yl, thiomorpholin-4-yl, piperidin-1-yl; and

heterocycles, except those of Ar and Ar^φ, can be substituted with in addition to any
5 substitutions specifically noted one or more substituents selected from acylamino, alkanoyl, alkoxy, alkoxycarbonyl, alkoxycarbonylalkyl, alkyl, (C₁ to C₃)alkylenedioxy, alkylamino, alkylsulfonyl, alkylsulfinyl, alkylthio, amino, ArC(O)-, ArO-, Ar-, Ar-alkyl, carboxy, dialkylamino, fluoro, fluoroalkyl, difluoroalkyl, hydroxy, mercapto, oxo, sulfamoyl, trifluoromethyl, 4-[C₆ or
10 C₁₀]arylpiperidin-1-yl and 4-[C₆ or C₁₀]arylpiperazin-1-yl;
or a pharmaceutically acceptable salt of said compounds,

with the proviso that where the compound of formula I is administered to decrease intraocular pressure at least one compound of formula I administered in effective amount is not a thiazole substituted on a ring carbon sulfonamide (the amide of
15 which can be substituted) that has carbonic anhydrase inhibiting activity.

2. The method of claim 1, comprising administering an intraocular pressure decreasing amount or ocular accommodation improving amount of a compound of the formula I, wherein the bond between carbons 4 and 5 is a single bond.

3. The method of claim 1, comprising administering an intraocular pressure decreasing amount or ocular accommodation improving amount of a compound of the formula I, wherein R^c is amino, amino(C₁-C₅)alkyl, or amino(C₆ or C₁₀)aryl, or wherein the amino of any of the three groups can be substituted with

(a) Ar;

(b) Ar-Z-, Ar-alkyl-Z-, Ar-Z-alkyl, Ar-amino-Z-, Ar-aminoalkyl-Z-, or Ar-oxyalkyl-Z-; or

(c) formyl or alkanoyl.

4. The method of claim 1, comprising administering an intraocular pressure decreasing amount or ocular accommodation improving amount of a compound of the formula I, wherein J is S or O, and R^c is hydrogen, oxo, alkyl, amino, amino(C₁-C₅)alkyl or aminophenyl, wherein the amino of the latter three groups can be substituted with

- (a) Ar;
- (b) Ar-Z-, Ar-alkyl-Z-, Ar-Z-alkyl, Ar-amino-Z-, Ar-aminoalkyl-Z-, or Ar-oxyalkyl-Z- ; or
- (c) formyl or alkanoyl.

5

5. The method of Claim 1, comprising administering an intraocular pressure decreasing amount or ocular accommodation improving amount of a compound of the formula I, wherein J is S, and R^c is hydrogen, oxo, alkyl, amino, amino(C₁-C₅)alkyl or aminophenyl, wherein the amino of the latter three groups can be substituted with

10

- (a) Ar;
- (b) Ar-Z-, Ar-alkyl-Z-, Ar-Z-alkyl, Ar-amino-Z-, Ar-aminoalkyl-Z-, or Ar-oxyalkyl-Z- ; or
- (c) formyl or alkanoyl.

15

6. The method of claim 1, comprising administering an intraocular pressure decreasing amount or ocular accommodation improving amount of a compound of the formula I, wherein the compound is selected from the group consisting of thiazole, 2-amino-4-chlorobenzothiazole, 2,4,5-trimethylthiazole, 2-(3,5-dimethylphenoxy)-N-thiazol-2-yl)acetamide, 2-isobutylthiazole, (4-fluorophenyl)thiazolin-2-ylamine, 2-furyl-N-[4-(6-methylbenzothiazol-2-yl)phenyl]carboxamide, and 5,5-dimethyl-2-(2-naphthylamino)-4,5,6-trihydrobenzothiazol-7-one.

20

7. The method of claim 1, comprising administering an intraocular pressure decreasing amount or ocular accommodation improving amount of a compound of the formula I, wherein

25

d. R^a and R^b are

30

1. independently selected from hydrogen, acylamino, alkanoyl, alkanoylalkyl, alkoxy, alkoxycarbonyl, alkoxycarbonylalkyl, alkyl, alkylamino, amino, ω-alkylenesulfonic acid, carbamoyl, carboxy, carboxyalkyl (which alkyl can be substituted with alkyloxyimino), cycloalkyl, dialkylamino, halo, hydroxy, (C₂-C₆)hydroxyalkyl, mercapto, nitro, sulfamoyl, sulfonic acid, alkylsulfonyl, alkylsulfinyl, alkylthio, trifluoromethyl, morpholin-4-yl, thiomorpholin-4-yl, piperidin-1-yl, 4-[C₆ or C₁₀]aryl piperidin-1-yl, 4-[C₆ or C₁₀]aryl piperazin-1-yl, Ar

- {wherein, consistent with the rules of aromaticity, Ar is C₆ or C₁₀ aryl or a 5- or 6-membered heteroaryl ring, wherein the 6-membered heteroaryl ring contains one to three atoms of N, and the 5-membered heteroaryl ring contains from one to three atoms of N or one atom of O or S and zero to two atoms of N, each
- 5 heteroaryl ring can be fused to a substituted benzene, pyridine, pyrimidine, pyrazine, pyridazine, or (1,2,3)triazine (wherein the ring fusion is at a carbon-carbon double bond of Ar)}, Ar-alkyl, ArO-, ArSO₂-, ArSO-, ArS-, ArSO₂NH-, ArNH-, (N-Ar)(N-alkyl)N-, ArC(O)-, ArC(O)NH-, ArNH-C(O)-, and (N-Ar)(N-alkyl)N-C(O)-; or
- 10 2. together with their ring carbons form a C₆- or C₁₀- aryl fused ring; or
3. together with their ring carbons form a C₅-C₇ fused cycloalkyl ring having no double bonds except any fused double bond of the formula I or IA ring, which cycloalkyl ring can be substituted by one or more of the group consisting of alkyl, amino, aminocarbonyl, carboxy, fluoro, or oxo, where multiple substituents are
- 15 located on different carbon atoms of the cycloalkyl ring, except in the case of alkyl and fluoro substituents, which can be located on the same or different carbon atoms; or
4. together with their ring carbons form a fused 5- or 6-membered heteroaryl ring, wherein the 6-membered heteroaryl ring contains one to three atoms of N, and
- 20 the 5-membered heteroaryl ring contains from one to three atoms of N or one atom of O or S and zero to two atoms of N; or
5. together with their ring carbons form a fused five to six membered second heterocycle, wherein the fused heterocycle consists of ring atoms selected from the group consisting of carbon, nitrogen, oxygen, sulfur, and S(O)_n, wherein n is
- 25 1 or 2,
- wherein aryl, Ar, or Ar^ϕ can be substituted with, in addition to any substitutions specifically noted one or more substituents selected from the group of alkyl, amino, dialkylamino, 1-pyrrolidinyl, 4-[C₆ or C₁₀]arylpiperazin-1-yl, 4-[C₆ or C₁₀]arylpiperidin-1-yl, azetidin-1-yl, morpholin-4-yl, thiomorpholin-4-yl,
- 30 piperidin-1-yl; and
- heterocycles, except those of Ar and Ar^ϕ, can be substituted with in addition to any substitutions specifically noted one or more substituents selected from acylamino,

5

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